

Unclassified

DTIC FULL COPY

2

SECURITY CLASSIFICATION OF THIS PAGE

## REPORT DOCUMENTATION PAGE

Form Approved  
OMB No 0704-0188

1a. REPORT SECURITY CLASSIFICATION		1b. RESTRICTIVE MARKINGS	
<b>AD-A230 034</b>		3. DISTRIBUTION/AVAILABILITY OF REPORT	
		Distribution unlimited	
		5. MONITORING ORGANIZATION REPORT NUMBER(S)	
Technical Report #11			
6a. NAME OF PERFORMING ORGANIZATION Massachusetts Institute of Technology	5b. OFFICE SYMBOL (If applicable)	7a. NAME OF MONITORING ORGANIZATION Office of Naval Research	
6c. ADDRESS (City, State, and ZIP Code) Department of Chemistry, 6-331 Cambridge, MA 02139		7b. ADDRESS (City, State, and ZIP Code) 800 North Quincy Street Arlington, VA 22217-5000	
8a. NAME OF FUNDING/SPONSORING ORGANIZATION Office of Naval Research	8b. OFFICE SYMBOL (If applicable)	9. PROCUREMENT INSTRUMENT IDENTIFICATION NUMBER	
8c. ADDRESS (City, State, and ZIP Code) 800 North Quincy Street Arlington, VA 22217-5000		10. SOURCE OF FUNDING NUMBERS	
		PROGRAM ELEMENT NO	PROJECT NO
		TASK NO	WORK UNIT ACCESSION NO.
11. TITLE (Include Security Classification) Polymerization of Functionalized Norbornenes Employing Mo(CH-t-Bu)(NAr)(O-t-Bu <sub>2</sub> ) as the Initiator.			
12. PERSONAL AUTHOR(S) G. C. Bazan, R. R. Schrock, H.-N. Cho and V. C. Gibson			
13a. TYPE OF REPORT Technical	13b. TIME COVERED FROM _____ TO _____	14. DATE OF REPORT (Year, Month, Day) December 5, 1990	15. PAGE COUNT
16. SUPPLEMENTARY NOTATION Submitted for publication in Macromolecules			
17. COSATI CODES		18. SUBJECT TERMS (Continue on reverse if necessary and identify by block number)	
FIELD	GROUP	alkylidene, molybdenum, metathesis polymerization, norbornenes	
19. ABSTRACT (Continue on reverse if necessary and identify by block number) Norbornenes containing esters, cyano, acetate, and related functionalities can be polymerized in a living manner employing Mo(CH-t-Bu)(NAr)(O-t-Bu) <sub>2</sub> as the initiator, especially in THF, to give polymers with polydispersities as low as 1.04. Living polymers containing Mo-alkylidene end groups have been characterized by proton NMR methods.			
20. DISTRIBUTION/AVAILABILITY OF ABSTRACT <input checked="" type="checkbox"/> UNCLASSIFIED/UNLIMITED <input type="checkbox"/> SAME AS RPT <input type="checkbox"/> DTIC USERS		21. ABSTRACT SECURITY CLASSIFICATION Unclassified	
22a. NAME OF RESPONSIBLE INDIVIDUAL Richard R. Schrock		22b. TELEPHONE (Include Area Code) 617/253-1596	22c. OFFICE SYMBOL

DD Form 1473, JUN 86

Previous editions are obsolete

SECURITY CLASSIFICATION OF THIS PAGE

S/N 0102-LF-014-6603

## DISTRIBUTION STATEMENT A

Approved for public release  
Distribution Unlimited

OFFICE OF NAVAL RESEARCH

Contract N00014-89-J1542

R&T Code 4132038

Technical Report No. 11

Polymerization of Functionalized Norbornenes Employing  
Mo(CH-*t*-Bu)(NAr)(O-*t*-Bu)<sub>2</sub> as the Initiator.  
by

G. C. Bazan,<sup>‡</sup> R. R. Schrock,<sup>\*†</sup> H.-N. Cho<sup>+</sup> and V. C. Gibson,<sup>§</sup>

Submitted for Publication

in

*Macromolecules*

<sup>‡</sup>California Institute of Technology  
Department of Chemistry  
1201 East California Boulevard  
Pasadena, California 91125 USA

<sup>†</sup>Massachusetts Institute of Technology  
Department of Chemistry, 6-331  
77 Massachusetts Avenue  
Cambridge, MA 02139 USA

<sup>+</sup>Korea Advanced Institute of Science and Technology  
Polymer Materials Lab  
Cheongryang, Seoul, KOREA

<sup>§</sup>University of Durham  
Department of Chemistry  
South Road, Durham DH1 3LE  
UNITED KINGDOM

December 5, 1990

Reproduction in whole or in part is permitted for any purpose of the United States Government.

This document has been approved for public release and sale; its distribution is unlimited.

Polymerization of Functionalized Norbornenes Employing  
 $\text{Mo}(\text{CH-}i\text{-Bu})(\text{NAr})(\text{O-}i\text{-Bu})_2$  as the Initiator.

by

G. C. Bazan, R. R. Schrock,\* H.-N. Cho and V. C. Gibson

Department of Chemistry 6-331  
 Massachusetts Institute of Technology  
 Cambridge, Massachusetts 02139

**Abstract**

Norbornenes containing esters, cyano, acetate, and related functionalities can be polymerized in a living manner employing  $\text{Mo}(\text{CH-}i\text{-Bu})(\text{NAr})(\text{O-}i\text{-Bu})_2$  as the initiator, especially in THF, to give polymers with polydispersities as low as 1.04. Poly-exo-cis-2,3-norbornenediacetate (poly-6), poly-exo-syn-2,7-norbornenediacetate (poly-7), and poly-exo-cis-2,3-norbornenediol-di-O-isopropylidene (poly-8) can be hydrolyzed to give polymers containing hydroxyl groups in place of acetates. Poly-6 and poly-7 lose two equivalents of acetic acid above 300 °C to give black intractable films; the analogous diols lose two equivalents of water under similar conditions. Addition of exo,cis-2,3-dichloro-endo,cis-2,3-carbonatonorbornene (**13**) to  $\text{Mo}(\text{CH-}i\text{-Bu})(\text{NAr})(\text{O-}i\text{-Bu})_2$  yields an isolable square pyramidal molybdacyclobutane complex, ~50% of which loses **13** at room temperature to reform  $\text{Mo}(\text{CH-}i\text{-Bu})(\text{NAr})(\text{O-}i\text{-Bu})_2$ . A block copolymer of 5-cyanonorbornene (**5**) and norbornene (100/100eq) was prepared and shown to be considerably stronger than a 200-mer of poly-5 (55% vs. 8% elongation before failure). Living polymers containing Mo-alkylidene end groups have been characterized by proton NMR methods.



By _____	
Distribution/	
Availability Codes	
Dist	Avail and/or Special
A-1	

## INTRODUCTION

Metathesis catalysts that will tolerate functionalities have been sought ever since the discovery of the olefin metathesis reaction.<sup>1</sup> Since the metal center in metathesis catalysts shows electrophilic character, there may be some direct correlation between the rate at which a metal-carbon double bond reacts with a carbon-carbon double bond and the rate at which it reacts with donor functionalities such as the carbonyl group. Therefore a given metal-carbon double bond could react relatively selectively with an especially reactive double bond (e.g., in a strained cyclic olefin) relative to a functionality elsewhere in the molecule. Indeed there seem to be many more examples of metathesis reactions (polymerizations) of norbornenes that contain functional groups remote from the double bond than ordinary olefins that contain remote functional groups.<sup>1</sup> However, since details concerning the structure and reactivity of classical metathesis catalysts are virtually nonexistent, their reactivities cannot be controlled systematically. In contrast a well-defined catalyst could be "deactivated" in a systematic fashion to the extent that it does not react with the functionality but still will react with the strained carbon-carbon bond on the timescale of the polymerization.

Recently well-characterized molybdenum<sup>2</sup> and tungsten<sup>3</sup> alkylidene complexes of the type  $M(\text{CH-}t\text{-Bu})(\text{NAr})(\text{OR})_2$  ( $\text{Ar} = 2,6\text{-diisopropylphenyl}$ ) have been synthesized whose reactivities toward carbon-carbon double bonds can be varied dramatically by varying the nature of the alkoxides. Complexes containing *t*-butoxides do not react readily with internal olefins, but will react readily with the double bond in norbornenes.<sup>4</sup> This results in a process where no chain termination or chain transfer occurs on the timescale of initiation and propagation, i.e., one that has all the characteristics of a living polymerization. Preliminary<sup>5</sup> and more recent<sup>6</sup> results have suggested that  $\text{Mo}(\text{CH-}t\text{-Bu})(\text{NAr})(\text{O-}t\text{-Bu})_2$  is much more tolerant of functional groups than  $\text{W}(\text{CH-}t\text{-Bu})(\text{NAr})(\text{O-}t\text{-Bu})_2$ . Furthermore intermediate alkylidene complexes that contain molybdenum seem to be much more stable toward decomposition and side reactions.<sup>7</sup> In this paper we report the polymerization of a variety of functionalized norbornenes by  $\text{Mo}(\text{CH-}t\text{-Bu})(\text{NAr})(\text{O-}t\text{-Bu})_2$ . The development of living ring-opening metathesis polymerization (ROMP<sup>8</sup>)

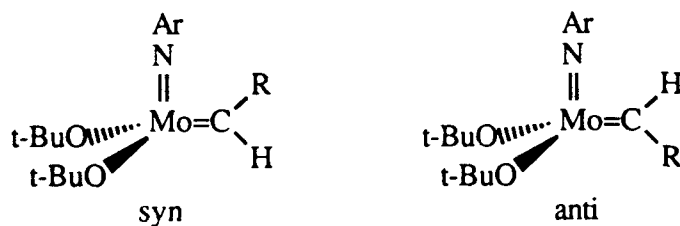
catalysts that tolerate functionalities should create the possibility of preparing a wide variety of functionalized polymers of this general type, and perhaps also create the possibility of attaining a significant degree of control over the polymer structure.<sup>6</sup>

## RESULTS

### Survey of functionalities tolerated by $\text{Mo}(\text{CH-}i\text{-Bu})(\text{NAr})(\text{O-}i\text{-Bu})_2$ .

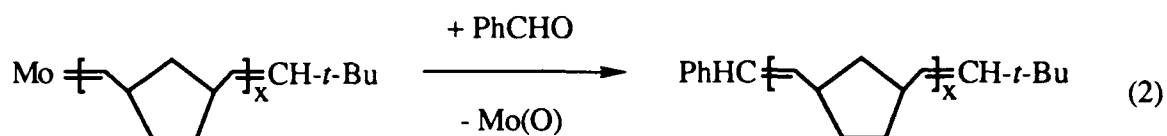
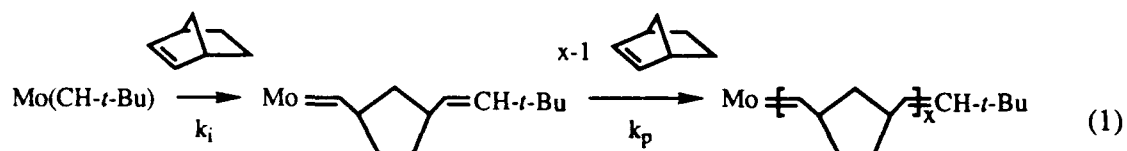
Monomers that have been successfully polymerized successfully by  $\text{Mo}(\text{CH-}i\text{-Bu})(\text{NAr})(\text{O-}i\text{-Bu})_2$  ( $\text{Mo}(\text{CH-}i\text{-Bu})$ ) are shown in Table I. A polymerization is termed successful if the monomer is consumed quantitatively according to proton NMR studies, the polymer can be cleaved from the metal in a Wittig-like reaction with aldehydes, and the polymer can be isolated quantitatively and has a narrow polydispersity (PDI). All of the polymers we have prepared are soluble in toluene or dichloromethane in the molecular weight range indicated. The preferred aldehydes that are used in the termination step do not have  $\beta$  protons (e.g., pivaldehyde or benzaldehyde). All polymerizations were carried out at room temperature under a nitrogen atmosphere in toluene or THF (~10 mL) with typical catalyst concentrations being 5 to 20 mM.

Norbornene (NBE) is polymerized extremely rapidly in toluene, benzene, or THF. The alkylidene resonance for  $\text{Mo}(\text{CH-}i\text{-Bu})$  at 11.23 ppm in  $\text{C}_6\text{D}_6$  is replaced by one new  $\text{H}_\alpha$  doublet resonance at 11.53 ppm that broadens slightly as the chain lengthens. There is no evidence for formation of a mixture of *syn* and *anti* rotamers about the  $\text{Mo}=\text{C}$  bond in living polynorbornene.<sup>2,3</sup>



By using 10 equiv of norbornene and measuring the amount of residual initiator the ratio  $k_p/k_i$  can be determined.<sup>6b</sup> The result ( $k_p/k_i = 12.1(5)$ )<sup>9</sup> is within the range that allows low polydispersities to be achieved for polymers containing 50 or more monomer units.<sup>10</sup> Olefinic proton resonances

characteristic of cis and trans double bonds in the polymer chain are observed in the ratio of 40:60.



Living polynorbornene reacts with benzaldehyde rapidly and quantitatively to yield benzylidene-capped polynorbornene (approximately 65 % trans configuration).

Comparison of the polymers from **4** (PDI=1.12, 100 equiv), **2** (PDI=1.07, 100 equiv), and **3** (PDI=1.04, 100 equiv) suggests that monomers that contain two endo functionalities behave less ideally than those that contain one or two exo substituents. In general polymerizations employing classical catalysts have been less successful if substituents (especially two) are in endo positions.<sup>1</sup> To our knowledge this is the first successful report of the polymerization of **2** and **4**. Polymerization of **3** by classical catalysts has been reported,<sup>1a</sup> but this polymerization, like the vast majority of those involving tungsten-based and molybdenum-based classical catalysts, did not show characteristics of a living system. Classical catalysts based on later transition metals such as ruthenium appear to tolerate functionalities to a greater degree<sup>1</sup> and recent studies have shown that functionality-tolerant ruthenium catalysts can be prepared that are active in water and that have some living characteristics.<sup>11</sup>

The proton NMR spectrum of poly-**2** suggests that propagation is ~95% trans selective, and that the trans to cis ratio does not change in living poly-**2** containing 20 equiv of monomer in several days, i.e., secondary metathesis is slow. The proton NMR spectrum of an oligomer made by the addition of 20 equiv of **2** to Mo(CH-*i*-Bu) followed by capping with benzaldehyde is shown in Figure 1. The configuration about the first double bond in the polymer is >98% trans (the error is ~2 %), as shown by the patterns for the resonances ascribable to H<sub>b</sub> at 5.29 ppm (dd, J<sub>HH</sub> = 15

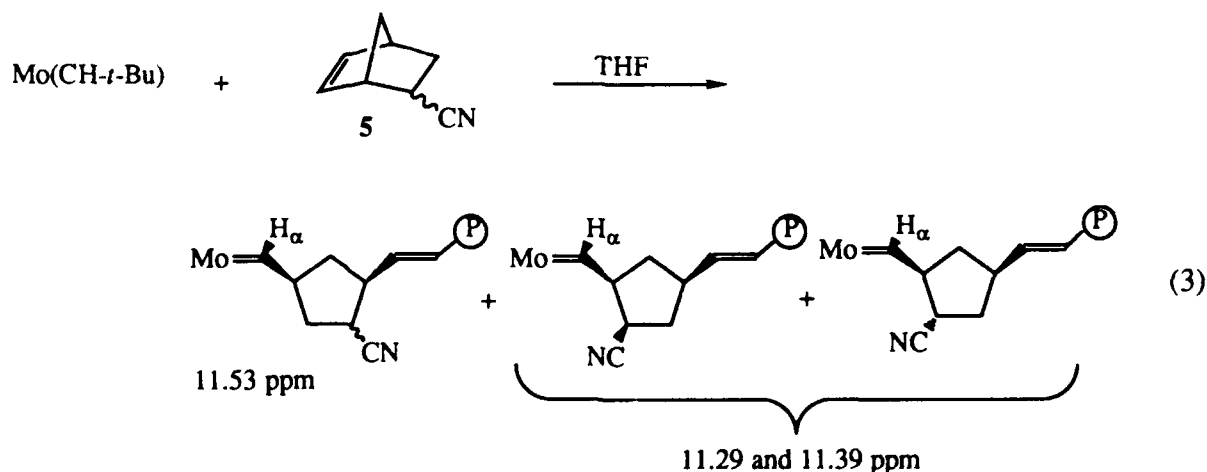
and 7 Hz) and  $H_a$  at 5.53 ppm (d,  $J_{HH} = 15$  Hz). The double bond that is formed in the Wittig-like reaction with benzaldehyde is also all trans with resonances at 6.02 ppm for  $H_c$  (dd,  $J_{HH} = 16$  and 7.5 Hz) and 6.42 ppm for  $H_d$  (d,  $J_{HH} = 16$  Hz). (The resonances at 6.18 and 6.32 ppm are olefinic resonances in *trans*-PhCH=CH-*t*-Bu that is formed upon reaction of residual Mo(CH-*t*-Bu) with benzaldehyde.) Presumably the greater bulk of the substituent in the alkylidene ligand forces an all-trans 2-oxamolybdacyclobutane intermediate to form.<sup>12</sup> Integration of the four olefinic protons on the two terminal double bonds versus the resonance for the *tert*-butyl capping group at 0.97 ppm shows the ratio to be approximately 1:1:1:1:9, consistent with quantitative yields in each of the proposed steps. If the polymerization is terminated by addition of pivaldehyde then only the resonances due to the olefinic protons near the *tert*-butyl end group (at 5.53 ppm 5.28 ppm) are observed. The proton and carbon NMR spectra can be assigned by a combination of homonuclear and heteronuclear correlation experiments. The two alkylidene proton resonances that are observed in the living oligomers at 11.51 and 11.45 are tentatively assigned to syn and anti rotamers (not necessarily respectively). (See Experimental Section for details.) Rotamers have been observed in polymers made from functionalized norbornadienes,<sup>6</sup> and there is no reason why they could not be present in these circumstances.

In proton NMR spectra of oligomers made from **4** the resonances due to cis olefinic protons overlaps with those due to trans olefinic protons but the cis:trans content can be estimated to be approximately 20:80 on the basis of the ratio of the allylic protons next to a cis or a trans olefinic bond. Resonances that could be ascribed to olefinic protons in the first and last double bond that arise in the initiation or termination steps cannot be discerned clearly.

Polymerization of **3** is expected to yield head-to-tail, tail-to-tail, and head-to-head sequences. Proton and carbon NMR spectra of the isolated polymers are broad and complex, consistent with a complex and random microstructure. Little information can be obtained. The alkylidene region of the proton NMR spectrum of living poly-**3** is shown in Figure 2. The two groups of alkylidene can be assigned to isomers in which the carbomethoxy functionality nearest the alkylidene carbon atom is either exo ( $H_\alpha$  at 11.55 ppm) or endo ( $H_\alpha$  at 11.65 ppm) by

comparison with alkylidene resonances in living poly-2 and living poly-4. Two resonances for the tert-butyl group from the initiator are observed at 0.964 and 0.935 ppm, depending on whether the carbomethoxy group is syn or anti relative to the trans-olefinic bond. Addition of 10 equiv of norbornene to living poly-3 produces an alkylidene resonance typical of living polynorbornene at 11.53 ppm. The two resonances for the tert-butyl group are unchanged, and furthermore, these solutions are stable for days; only the alkylidene resonance for living polynorbornene is observed with its original intensity.

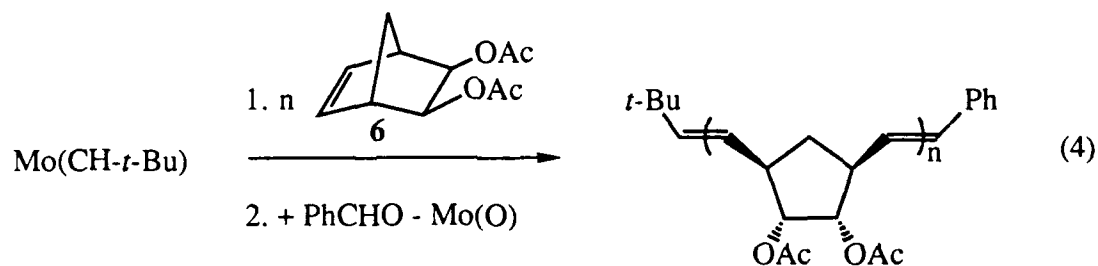
Addition of 100-200 equiv of 5-cyanonorbornene (**5**) to  $\text{Mo}(\text{CH-}i\text{-Bu})$  in toluene does not yield polymer. However, in THF as many as 200 equiv of **5** are polymerized smoothly to give polymers with polydispersities of 1.06-1.07. The alkylidene proton region of the proton NMR spectrum in THF- $d_8$  for living poly-5 is shown in Figure 3. The three main groups of resonances that are observed can be rationalized in terms of the three basic isomers that are possible upon opening this unsymmetric monomer (eq 3). The resonance at 11.53 ppm is assigned to the alkylidene resonance which most resembles that in living polynorbornene (eq 3). We can only speculate as to the source of the doubling of this resonance. Possibilities are that the chemical shift of  $\text{H}_\alpha$  is sensitive to exo or endo orientation of the cyano group relative to the  $\text{C}_\alpha\text{-C}_\beta$  bond or cis or trans configuration of the first  $\text{C}=\text{C}$  bond, or syn and anti rotamers are present. Whatever the explanation the ratio of the two  $\text{H}_\alpha$  types is approximately 1:1. The two other resonances at





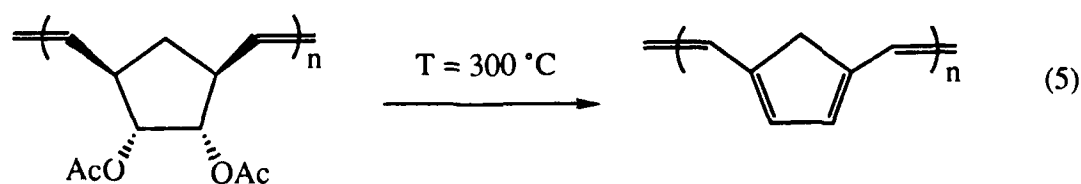
$\sim 11.29$  and  $\sim 11.39$  ppm are assigned to  $H_\alpha$  protons in propagating alkylidene complexes in which the cyano group (either exo or endo) is on a  $\gamma$  carbon atom with respect to the metal. In this case the ratio is not 1:1. Further complexity within each group of resonances could arise from a sensitivity to the nature of adjacent monomers in the chain as well as from the presence of syn and anti rotamers. The identification of these resonances is further supported by the observation of only two complex envelopes of resonances in the living polymer prepared from 5,6-transdicyanonorbornene.<sup>13</sup>

**6** is polymerized quantitatively in THF to give a polymer with a polydispersity of 1.19 (100-mer). In aromatic solvents (toluene, chlorobenzene, or benzene) **6** is not consumed



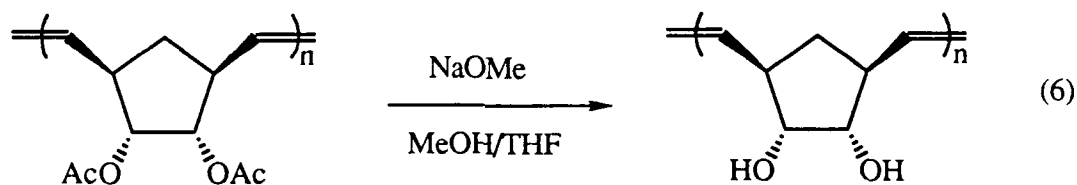
completely ( $\sim 85\%$ ) and broader molecular weight distributions are found ( $\text{PDI} = 1.3\text{--}2$ ). The IR spectrum of the polymer shows the characteristic ester carbonyl absorption at  $1740 \text{ cm}^{-1}$ . By proton NMR the olefinic trans to cis ratio was determined to be 60:40.

Poly-**6** is a white powder which is insoluble in methanol and can be cast from toluene as a flexible transparent film. The tensile strength of the film at the breaking point is  $370 \text{ kg/cm}^2$ , a value that is comparable to that of acrylonitrile/butadiene/styrene (ABS) resin ( $380 \text{ kg/cm}^2$ ).<sup>14</sup> A DSC measurement showed only a glass transition temperature ( $T_g$ ) at  $102^\circ\text{C}$ . TGA suggests that



the polymer is stable up to 300 °C. Above 300 °C two equiv of acetic acid are lost (Fig 4, eq 5) followed by polymer decomposition at higher temperatures. Isothermal pyrolysis showed that at 300 °C 55.7 % of the mass is lost in 2 h while at 350 °C 60.9 % is lost in 1 h. (Loss of two equiv of acetic acid corresponds to a 57.2 % mass loss.) Heating a strip of polymer at 300 °C for two hours in a temperature programmable furnace yielded a red-black insoluble film. Its IR spectrum showed that the carbonyl absorption characteristic of the precursor polymer has decreased and the 1600  $\text{cm}^{-1}$  (C=C) absorption has increased. Therefore pyrolysis appears to be incomplete. When the black polymer is left exposed to air an absorption band at 1730  $\text{cm}^{-1}$  appears, characteristic of some oxidation process, the nature of which remains undetermined.

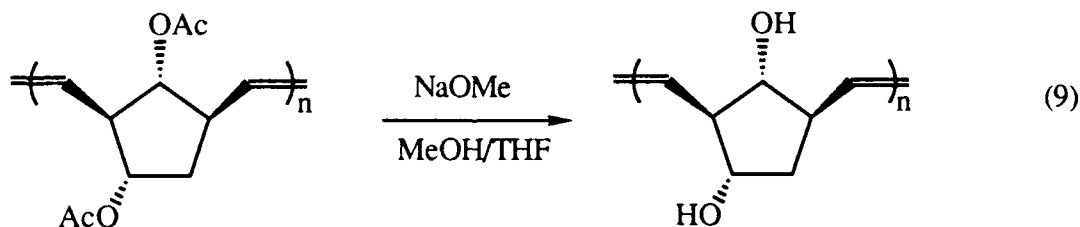
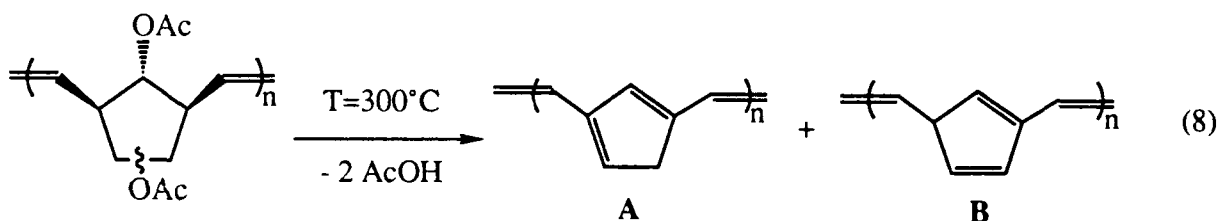
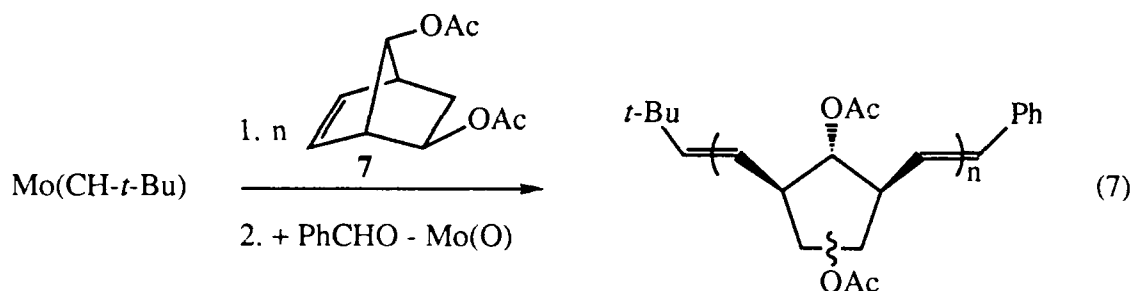
Hydrolysis of poly-6 gives the polydiol shown in equation 6 quantitatively. Since hydroxyl groups are incompatible with  $\text{Mo}(\text{CH-}t\text{-Bu})(\text{NAr})(\text{O-}t\text{-Bu})_2$  this polymer cannot be made directly from the corresponding monomer. The polydiol is insoluble in standard solvents



(THF, chloroform, aromatics, methanol, or water) but it dissolves completely in acidic chloroform ( $\text{CF}_3\text{CO}_2\text{H}/\text{CDCl}_3$ ). The solid polymer absorbs strongly in the region between 3100 and 3500  $\text{cm}^{-1}$ ; the absorption at 1740  $\text{cm}^{-1}$  in the precursor polymer is absent. No glass transition was found in the polydiol by DSC up to 220 °C. The TGA displayed a two stage degradation, the first step of which (at ~300 °C) corresponds to a dehydration analogous to the thermal elimination shown in equation 5.

Exo-syn-5-norbornene-2,7-diol diacetate (7; eq 7) is polymerized quantitatively to yield a polymer with a PDI = 1.09 in THF and 2.17 in chlorobenzene (200 mer). Poly-7 can be purified by precipitation into methanol and can be cast from toluene to give flexible, transparent thin films. Its tensile strength at the breaking point is 420  $\text{kg}/\text{cm}^2$ . DSC analysis indicates that it is amorphous

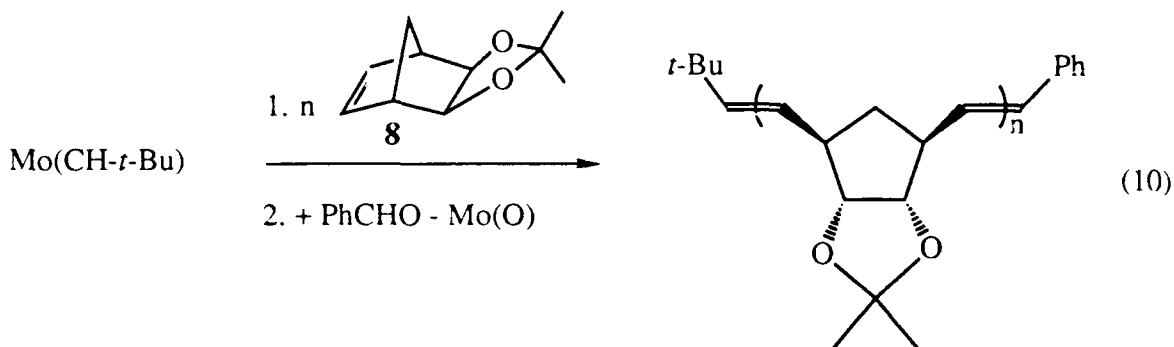
with a  $T_g$  at 110 °C and that it is stable up to 300 °C. The TGA for poly-7 resembles that for poly-6, consistent with loss of two equiv of acetic acid from the polymer backbone (eq 8). Structure A is likely to be more favorable as a result of cross conjugation. The pyrolysis product made from poly-7 is brittle and oxidizes readily in air.



Poly-7 can be hydrolyzed to yield the polydiol shown in equation 9. The IR spectrum of poly-7 shows a broad OH absorption at 3100-3500  $\text{cm}^{-1}$ , a vinyl stretch at 1590  $\text{cm}^{-1}$ , but no carbonyl absorption. A proton NMR spectrum in a 1:1 mixture of  $\text{CF}_3\text{CO}_2\text{H}$  and  $\text{CDCl}_3$  showed the hydroxyl protons as a broad resonance at  $\delta$  2.8 ppm. This polymer also is soluble in *aqueous* acidic media ( $\text{CF}_3\text{CO}_2\text{H}/\text{H}_2\text{O}$ ). The TGA is similar to that of the polydiol shown in equation 6, the first stage of which can be attributed to loss of two equiv of water to give the polymers shown in equation 8, most likely with structure A. Isothermal pyrolysis at 270°C shows the loss of mass

to be 25.8% in 2 h (theoretical mass loss for dehydration = 28.6%).

**8** (eq 10) can be polymerized in THF straightforwardly. Proton NMR spectra of poly-**8** shows it to contain 60 % trans olefinic linkages. TGA analysis reveals a Tg at 108 °C and decomposition starting at 300 °C. Hydrolysis of poly-**5** yields the polydiol obtained by hydrolysis of poly-**6** mentioned earlier.

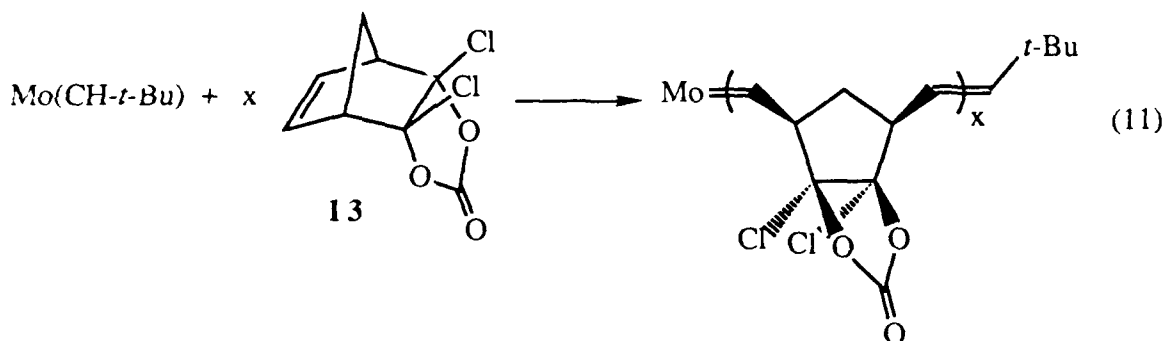


Monomers **9**, **10**, **11**, and **12** were polymerized smoothly to give the homopolymers shown in Table I, but the polymer properties were not investigated. It is interesting to note that the endo isomer analogous to **10** could not be polymerized by  $\text{Mo}(\text{CH-}t\text{-Bu})(\text{NAr})(\text{O-}t\text{-Bu})_2$ . As mentioned earlier, this type of result has been ascribed to steric problems associated with approach of an endo-substituted monomer to an alkylidene that has just been formed from that monomer.<sup>1</sup>

#### Reversible Formation of a Square Pyramidal Molybdacyle with exo,cis-2,3-dichloro-endo,cis-2,3-carbonatonorbornene.

Researchers have been interested in polymerizing exo,cis-2,3-dichloro-endo,cis-2,3-carbonato-norbornene (endo-3a,7a-dichloro-3a,4,7,7a-tetrahydro-4,7-methano-1,3-benzodioxol-2-one) (**13**) because poly-**13** is a potential precursor to conjugated materials. **13** has been polymerized using classical catalysts, but the polydispersity was large (76<sup>15,16</sup>). Reaction of 10 equiv of **13** with  $\text{Mo}(\text{CH-}t\text{-Bu})$  in  $\text{C}_6\text{D}_6$  at room temperature for approximately 60 min yields living oligomers (eq 11), judging from the broad alkylidene doublet at 11.57 ppm, but unreacted **13** remained. The monomer was completely consumed in 24 h at room temperature. The intensity of the alkylidene resonance for the living species appears to decrease slowly with time and attempts

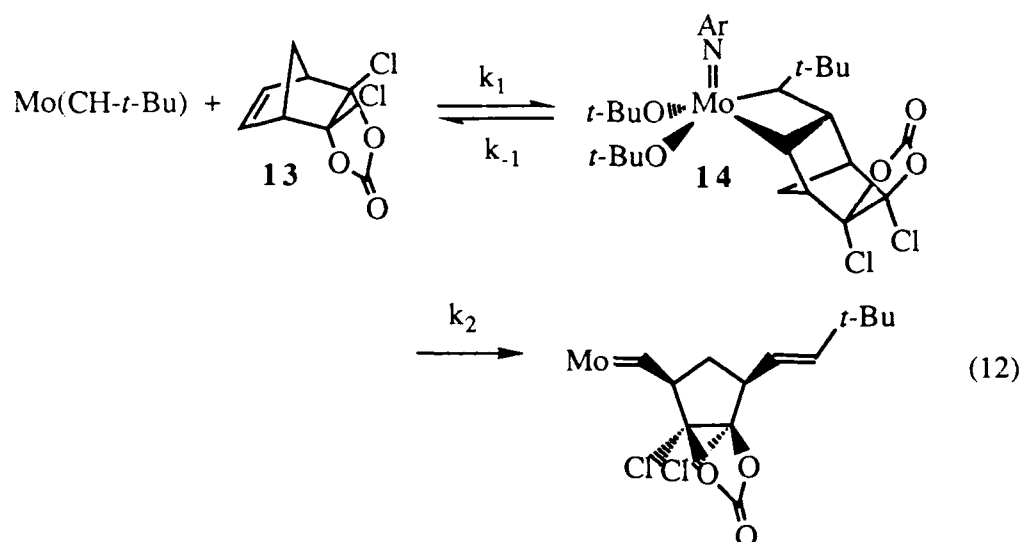
to polymerize 100 equiv of **13** in toluene in the standard fashion failed, both of which are believed to be the consequence of some decomposition of the living species. Increasing the reaction time at a catalyst concentration of 0.02 M up to 24 h at room temperature also did not yield any polymer.



Closer examination of the polymerization reaction by proton NMR showed that only approximately two equiv of **13** were consumed over a 2 hour period ( $[\text{Mo}(\text{CH-}t\text{-Bu})] = 0.02 \text{ M}$ , 4 equiv **13**, 25 °C). At -30 °C the neopentylidene and olefin resonances decreased but no new alkylidene resonances were found. However, broad resonances appeared which resembled those of well-characterized square pyramidal tungstacyclobutane complexes.<sup>17</sup> Therefore we propose that the square pyramidal molybdacycle (**14**; eq 12) forms at this temperature and does not open readily.  $\text{Mo}(\text{CH-}t\text{-Bu})$  was not completely converted to **14** in the presence of **13** at -30 °C.

A solution containing  $\text{Mo}(\text{CH-}t\text{-Bu})$  and **13** was prepared at -30°C and stored at -40 °C for 2-3 days. The bright red crystals that formed were dissolved in toluene- $d_8$  at -45 °C. At that temperature a mixture of 4 parts of **14** and one part of a 1:1 mixture of  $\text{Mo}(\text{CH-}t\text{-Bu})$  and **13** was observed. Only a trace of the propagating alkylidene was observed (Table II). As the temperature was raised the amount of propagating alkylidene,  $\text{Mo}(\text{CH-}t\text{-Bu})$  and **13** increased relative to **14**. At 0 °C the ratio of **14** to  $\text{Mo}(\text{CH-}t\text{-Bu})$  was 2:1. Finally at 25 °C only traces of **14** were observed and **13** was regenerated to give approximately 60 %  $\text{Mo}(\text{CH-}t\text{-Bu})$ . These results indicate that **14** is unstable at 25 °C but it "decomposes" either by ring-opening to yield a new propagating alkylidene or degeneratively to yield the original reactants (eq 12), i.e.,  $k_1[\mathbf{14}] \approx k_2$ . At low temperatures **14** is in equilibrium with  $\text{Mo}(\text{CH-}t\text{-Bu})$  and **13** ( $K = k_1/k_{-1} = 4$  at -45 °C) while

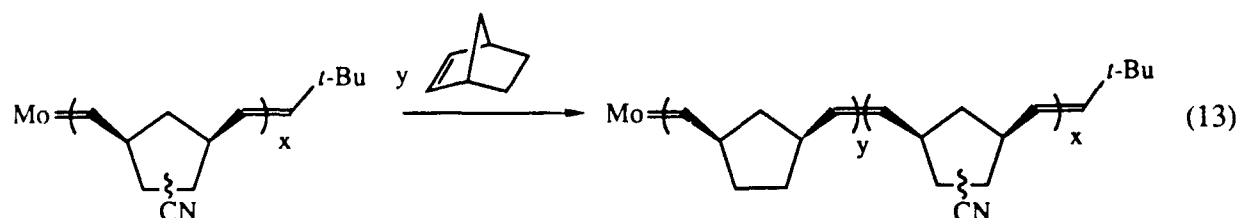
formation of new alkylidene is negligible ( $k_2$  is small). At 0 °C  $K \approx 2$ .



Failure to polymerize **13** can be attributed to the slow rate of its reaction with either  $\text{Mo}(\text{CH-}t\text{-Bu})$  or the propagating alkylidene complex. In this case the rate of catalyst deactivation, most likely in a reaction involving the carbonyl bond in the carbonate, is approximately the same as the rate of polymerization.

### Block Copolymers

Analysis by proton NMR shows that 15 equiv of norbornene are consumed upon addition to a solution of living poly-**5** and the complex set of alkylidene resonances characteristic of poly-**5** (Figure 3) are replaced by the doublet characteristic of living polynorbornene. This implies that poly-**5** reacts with norbornene to give a living block copolymer (eq 13). Addition of **5** to living polynorbornene yields the resonances ascribed to poly-**5**. If  $x = y = 100$  and the block copolymer is cleaved from the metal by addition of benzaldehyde, then a block copolymer (poly-1/5) whose GPC is shown in Figure 5 is obtained. Inverting the order of addition of the two monomers



produces a virtually identical polymer (poly-5/1) provided that more time is allowed for polymerization of **5**. Note that the PDI = 1.05 and peak is symmetric, indicative of a process in which there is no chain transfer or termination on the time scale of either polymerization.

The proton NMR spectrum of poly-1/5 is essentially a superposition of those of polynorbornene and poly-5 homopolymers. The olefinic region of the carbon NMR spectrum of poly-1/5 is compared with that of poly-5 in Figure 6. The complexity of the olefinic resonances in poly-5 can be attributed to the many variations of monomer sequences for this material. The resonances are broad since the olefinic protons are likely to be sensitive to the stereochemistry of the monomers on both sides of the olefinic bond. In poly-1/5 the resonances at 133.8 (cis olefin) and 132.9 (trans olefin) ppm can be ascribed to the polynorbornene block.

The stress-strain responses for poly-1/5 and poly-5 are shown in Figure 7. The behavior of poly-5 ( $T_g = 123\text{ }^\circ\text{C}$ ) is typical of what one would expect for either a glassy polymer or a semicrystalline polymer at a temperature below  $T_g$ ; the polymer collapses catastrophically at 7.9 % strain. Poly-1/5 is dramatically tougher, the break point occurring at 54.5 % strain.

## DISCUSSION

Narrow dispersities and the formation of block copolymers indicate that norbornenes containing the variety of functionalities shown in Table I are polymerized in a living manner by  $\text{Mo}(\text{CH}-t\text{-Bu})$ . For example the preparation of a 200 mer of **3** suggests that the catalytic site is stable in the presence of 400 equiv of ester functionality over the time period of the polymerization. Solvent plays an important role, although the precise nature of that role remains to be determined. One possibility is that THF competes successfully with the functionality for the metal. Another possibility is a bulk solvent effect that keeps the polar groups pointed away from the metal into the solution. A third possibility is that THF is intimately involved in opening the intermediate metallacycles formed in these reactions.

Liberation of **13** from **14** represents the first example of the reversible formation of a metallacyclobutane ring involving a norbornene derivative. This situation is likely to be rare and probably encountered only when the monomer is bulky and relatively electron poor. The high

stability of **14** suggests that a considerable amount of intramolecular rearrangement is necessary in order to lose the olefin from this metallacycle in a productive fashion. Similar conclusions have been reached recently in related circumstances.<sup>17a,6b</sup>

Ejection of small molecules in a precursor polymer to yield a conjugated polymer is a fairly mature subject in polymer chemistry.<sup>18</sup> Poly(vinyl acetate),<sup>19</sup> poly(vinyl chloride)<sup>20</sup> and poly(vinyl alcohol)<sup>21</sup> lose acetic acid, HCl and water respectively to yield conjugated polyene sequences at high temperatures. But pyrolysis of poly-**6** and poly-**7** and the analogous diols appear to be the first reports of such a reaction for polymers prepared by ring-opening metathesis. Most likely this circumstance can be ascribed to the previous paucity of catalysts for preparing suitably functionalized polymers.

Transformation of poly-**6**, poly-**7**, and poly-**8** to polydiols is interesting because of the unusual solubility properties of these materials and unobservable (presumably high)  $T_g$ . Poly(exo-5-hydroxynorbornene)<sup>22</sup> has been prepared by polymerizing an organoborane derivative of norbornene using classical catalysts followed by alkaline oxidation of the resulting polymer. The final polymer displays bulk characteristics similar to the ones reported here but the polydispersity was 2.3 and only 65 % of the monomer was consumed. An unsuccessful attempt to dehydrate the polymer was attributed to the unfavorable introduction of ring strain.

The change in mechanical properties observed for poly-**1/5** relative to poly-**5** is a typical example of the degree of control of the physical properties that is possible when the primary structure of the polymer can be controlled closely. The small strain for poly-**5** is typical of a glassy polymer (no melting temperature was observed by DSC). In poly-**1/5** the stress at maximum load is adequately maintained (and under these conditions occurs at approximately the same strain) but the break point of the material occurs at a much larger strain. Polymers which can maintain a high load and absorb energy upon fracture is a desirable property of materials for many applications.

## CONCLUSION

We have shown that several functionalized norbornenes can be polymerized in a living manner by  $\text{Mo}(\text{CH-}i\text{-Bu})(\text{NAr})(\text{O-}i\text{-Bu})_2$ , especially in THF, and that polymerization proceeds via



molybdacyclobutane and alkylidene intermediates. Success can be attributed to deactivation of the electrophilic metal toward metathesis of ordinary olefins by *t*-butoxide ligands, to the low electrophilicity of Mo (vs. W), to the use of THF as the solvent, and to the relatively high reactivity of the norbornene double bond. These results suggest that norbornenes that contain a variety of other functionalities (however, none that contain active protons) may be polymerized in a controlled manner under these conditions.

## EXPERIMENTAL SECTION

**General Details.** All polymerizations were performed under a nitrogen atmosphere in a Vacuum Atmospheres drybox or by using standard Schlenk techniques. Tetrahydrofuran was predried over calcium chloride and distilled from sodium benzophenone ketyl under nitrogen immediately prior to usage. Toluene was distilled from molten sodium and stored over sodium potassium alloy. All deuterated NMR solvents were passed through a column of activated alumina. Norbornene was doubly distilled from molten sodium. Commercially available **5** (60:40 mixture of *exo* and *endo* isomers) was distilled under nitrogen, passed through a column of activated alumina and stored under nitrogen. **12** is commercially available and **4** was prepared by adding methanol to the commercially available *endo*-anhydride.<sup>23</sup> Commercially available aldehydes were purified by distillation under nitrogen. **2**,<sup>11,24</sup> **3**,<sup>25</sup> **13**,<sup>14</sup> **6**,<sup>26</sup> **7**,<sup>27</sup> and **8**<sup>28</sup> were prepared by literature procedures.

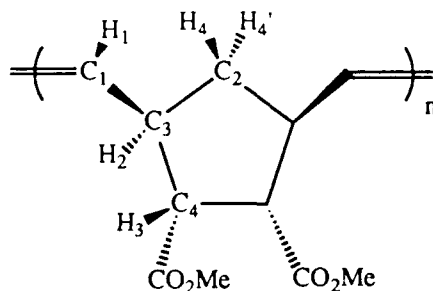
NMR data are listed in parts per million downfield from TMS for both proton and carbon. Coupling constants are quoted in Hertz. Obvious multiplicities and routine coupling constants usually are not listed. Gel Permeation Chromatographic (GPC) analysis were carried out at room temperature employing a Rheodyne Model 7125 sample injector, a Kratos Spectroflow 400 pump, Shodex KF-802.5, 803, 804, 805, 800P columns, a Knauer differential refractometer, and a Spectroflow 757 absorbance detector on samples 0.1-0.3 % w/v in dichloromethane which were filtered through a Millex-SR 0.5  $\mu$ m filter in order to remove particulates. GPC columns were calibrated versus commercially available polystyrene standards (Polymer Laboratories Ltd.) ranging from 1206 to  $1.03 \times 10^6$  MW. Thermal analysis (TGA) was done on a Perkin-Elmer

TGS-2 thermogravimetric analyzer. Differential scanning calorimetry was performed on a Perkin-Elmer instrument at a heating rate of 30 °C/min. Tensile strengths of polymer films (0.35 mm × 3mm : thickness × width) were measured by use of an Instron 4201 at an extension rate of 10 mm/min.

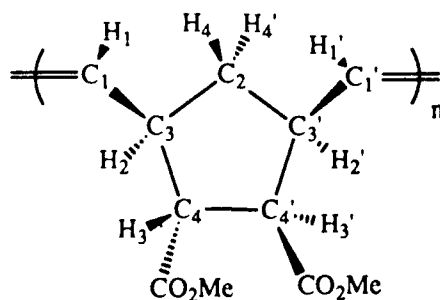
**Typical Polymerization of *exo,cis*-2,3-dicarbomethoxynorbornene (2).** A solution of **2** (431 mg, 2.05 mmol, 100 equiv) in toluene (10 mL) was added dropwise quickly to a rapidly stirring solution of Mo(CH-*t*-Bu)(NAr)(O-*t*-Bu)<sub>2</sub> (10 mg, 0.0205 mmol) in toluene (5 mL) over a period of 1-2 min. The solution was allowed to stir for an additional 20-30 min, quenched by addition of excess benzaldehyde (10 µL, ~ 0.1 mmol) and stirred for an additional 20 minutes. The solution was brought out of the box and added dropwise to 500 mL of vigorously stirred hexane. The resulting precipitate was isolated by centrifugation, washed with hexane or pentane, and placed under vacuum overnight. The final product was a white powder (380 mg, 88%) soluble in toluene, CH<sub>2</sub>Cl<sub>2</sub>, or benzene: <sup>1</sup>H NMR (CDCl<sub>3</sub>) δ 5.37 (br, 2, H<sub>1</sub>), 3.62 (s, 6, CO<sub>2</sub>Me), 2.93 (br, 2, H<sub>2</sub>), 2.79 (br, 2, H<sub>3</sub>), 2.06 (br, 1, H<sub>4</sub> or H<sub>4'</sub>), 1.25 (br, 1, H<sub>4</sub> or H<sub>4'</sub>), 0.96 (s, *t*-Bu); <sup>13</sup>C NMR δ 173.1 (CO<sub>2</sub>Me), 132.0 (C<sub>1</sub>), 52.4 (C<sub>4</sub>), 51.7 (CO<sub>2</sub>Me), 45.1 (C<sub>3</sub>), 39.1 (C<sub>2</sub>), 29.5 (*t*-Bu). GPC data are found in Table I.

Living oligomers were prepared by adding the monomer (65 mg, 0.31 mmol, 15 equiv) in C<sub>6</sub>D<sub>6</sub> (300 µL) to a rapidly stirred solution of Mo(CH-*t*-Bu)(NAr)(O-*t*-Bu)<sub>2</sub> (10 mg, 0.0205 mmol) in C<sub>6</sub>D<sub>6</sub>: <sup>1</sup>H NMR δ 11.51 and 11.45 (H<sub>α</sub>, J<sub>H<sub>α</sub>H<sub>β</sub></sub> = 7 Hz), 4.61 (H<sub>β</sub>, J<sub>H<sub>α</sub>H<sub>β</sub></sub> = 7 Hz), 3.97 (CHMe<sub>2</sub>). Resonances due to the oligomer protons are similar to those of the polymer; other resonances due to the ligands on the metal center could not be assigned due to overlap with oligomer resonances. Assignment of resonances from H<sub>2</sub>, H<sub>3</sub> and H<sub>4</sub> along with C<sub>2</sub>, C<sub>3</sub> and C<sub>4</sub> is ambiguous on chemical shift arguments alone. H<sub>2</sub> is assigned to the resonance at 2.93 ppm since it couples with H<sub>1</sub> (5.37 ppm). C<sub>2</sub> (39.1 ppm) is split into a triplet in a gated <sup>13</sup>C experiment which allows the assignment of H<sub>4</sub> and H<sub>4'</sub> (J<sub>C<sub>2</sub>H<sub>4</sub></sub> = J<sub>C<sub>2</sub>H<sub>4'</sub></sub>) to the resonances at 1.25 and 2.06 ppm since both are shown to be coupled in a HETCOR (<sup>13</sup>C-<sup>1</sup>H correlation) experiment to C<sub>2</sub>. The same experiment shows that the resonance at 45.1 ppm is coupled to H<sub>2</sub> and therefore belongs

to C<sub>3</sub>. By elimination, C<sub>4</sub> (52.4 ppm) and H<sub>3</sub> (2.79 ppm) can be assigned.



**Typical Polymerization of trans-2,3-dicarbomethoxynorbornene (3).** A solution of **3** (431 mg, 2.05 mmol, 100 equiv) in toluene (10 mL) was added dropwise but quickly to a rapidly stirring solution of Mo(CH-*t*-Bu)(NAr)(O-*t*-Bu)<sub>2</sub> (10 mg, 0.0205 mmol) in toluene (5 mL) over a period of 1-2 minutes. The solution was allowed to stir for an additional 20-30 minutes, quenched by addition of excess benzaldehyde (10  $\mu$ L, appx 0.1 mmol) (or pivaldehyde) and stirred for an additional 20 minutes. The solution was brought out of the box and added dropwise to 500 mL of hexane under vigorous stirring, the resulting precipitate was isolated by centrifugation, washed with hexane or pentane and placed under vacuum overnight. The polymer was obtained as white flakes (405 mg, 94%); <sup>1</sup>H NMR  $\delta$  (CDCl<sub>3</sub>, 500 MHz) 5.08 and 5.25 (br, 2, H<sub>1</sub> and H<sub>1'</sub>), 3.43 (s, 3, CO<sub>2</sub>Me), 3.40 (s, 3, CO<sub>2</sub>Me), 3.00 (br, 1, H<sub>2</sub> or H<sub>2'</sub> or H<sub>3</sub> or H<sub>3'</sub>), 2.73 (br, 2, H<sub>2</sub> or H<sub>2'</sub> or H<sub>3</sub> or H<sub>3'</sub>), 2.46 (br, 1, H<sub>2</sub> or H<sub>2'</sub> or H<sub>3</sub> or H<sub>3'</sub>), 1.76 (br, 1, H<sub>4</sub> or H<sub>4'</sub>), 1.27 (br, 1, H<sub>4</sub> or H<sub>4'</sub>), 0.935 and 0.962 (s, CMe<sub>3</sub>); <sup>13</sup>C 174.3 (CO<sub>2</sub>Me), 173.5 (CO<sub>2</sub>Me), 132.2 (C<sub>1</sub>), 131.2 (C<sub>1</sub>), 129-131.2 (C<sub>1</sub>), 51-53 (C<sub>4</sub> and CO<sub>2</sub>Me), 44.5 (C<sub>3</sub> or C<sub>3'</sub>), 46.7 (C<sub>3</sub> or C<sub>3'</sub>), 29.5 (CMe<sub>3</sub>). All resonances are extremely broad consistent with the many stereochemical configurations possible. GPC data are found in Table I.



Living oligomers were prepared by addition of the monomer (65 mg, 0.31 mmol, 15 equiv) in  $C_6D_6$  (300  $\mu$ L) to a rapidly stirring solution of  $Mo(CH-t-Bu)(NAr)(O-t-Bu)_2$  (10 mg, 0.0205 mmol) in  $C_6D_6$ . Oligomer resonances are similar to the polymer, for resonances due to  $H_\alpha$  see Figure 2.

**Polymerization of *exo-cis*-(*N*-phenyl)-5-norborne-2,3-dicarboximide (10).** A solution of **10** (239 mg, 1.03 mmol, 50 equiv) in THF (15 mL) was added dropwise over a period of 2-3 minutes to a rapidly stirring solution of  $Mo(CH-t-Bu)$  (10 mg) in THF (4 mL) and stirred an additional 30 minutes. The polymer was quenched with benzaldehyde and isolated as described above. The product was obtained as a gummy solid (213 mg, 90 %):  $^1H$  NMR  $\delta$  ( $CDCl_3$ ) 6.9-7.2 (br, 5,  $H_{Ar}$ ), 5.55 (br, 2,  $CH=CH$ ), 2.87 (br, 2,  $CHCO$  or allylic  $CH$ ), 2.55 (br, 2,  $CHCO$  or allylic  $CH$ ), 1.90 (br, 1,  $CHH$ ), 1.41 (br, 1,  $CHH$ ). GPC data:  $M_n=21000$  (vs polystyrene), PDI = 1.06.

**Polymerization of *endo-cis*(*N*-pentafluorophenyl)-5-norborne-2,3-dicarboximide (9).** Prepared as described above for the protio analog using 329 mg of monomer. The product was obtained as a white powder:  $^1H$  NMR  $\delta$  5.61 (br, 2,  $CH=CH$ ), 2.6-2.8 (br, 4,  $CHO$  and allylic  $CH$ ), 1.82 (br, 1,  $CHH$ ), 1.30 (br, 1,  $CHH$ ).

**Preparation of 14 and observation of its decomposition.** A solution of **13** (68 mg, 0.31 mmol) in ether (2 mL,  $-30^\circ C$ ) was layered over a solution of  $Mo(CH-t-Bu)$  (150 mg, 0.31 mmol) in pentane (2.0 mL,  $-30^\circ C$ ) and immediately stored at  $-40^\circ C$  for 48 hours. Bright red crystals (98 mg, 45%) were isolated by decanting the solution and briefly placed under vacuum. Analysis by  $^1H$  NMR was done by dissolving 20 mg of the product in toluene- $d_8$  (800  $\mu$ L, [**14**] =  $3.5 \times 10^{-5}$  M) at low temperature:  $^1H$  NMR ( $-45^\circ C$ ) 7.02 (m, 3,  $H_{Ar}$ ), 4.02 (br, 1,  $CHMe_2$ ), 3.83 (br, 1,  $CHMe_2$ ), 3.55 (s, 1, bridgehead H), 3.41 (dd, 1,  $H_\beta$ ,  $J_{H\alpha'H\beta}=8$ ,  $J_{H\alpha'H\beta}=8$ ), 2.62 (s, 1, bridgehead H), 1.85 (d, 1,  $CHH$  bridgehead ( $H_\gamma$ )), 1.68 (d, 1,  $CHH$  bridgehead ( $H_\gamma$ )), 1.32 (br, 12,  $CHMe_2$ ), 1.29 (s, 9,  $OCMe_3$ ), 1.17 (s, 9,  $OCMe_3$ ), 0.92 (s, 9,  $CHCMe_3$ ).

**Typical polymerization of *exo-cis*-2,3-norbornenediacetate (6).** A solution of **6** (431 mg, 2.05 mmol) in THF (10 mL) was added dropwise but quickly to a vigorously stirring

solution of Mo(CH-*t*-Bu) (10 mg,  $2. \times 10^{-5}$  mol) in THF (5 mL) over a period of 1-2 minutes. The resulting solution was allowed to stir for 1 hour, quenched with benzaldehyde (15  $\mu$ L, approx 0.15 mmol) and allowed to stir for an additional 30 minutes. The polymer was purified by dropwise addition of the resulting solution into a large excess of methanol, filtration, a subsequent precipitation from chloroform into methanol and placing under vacuum for 24 hours. The yield of isolated polymer (410 mg, 95 %) was essentially quantitative:  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  5.42 (br, CH=CH, trans), 5.33 (br, CH=CH, cis), 4.96 (br, CHOAc), 4.87 (br, CHOAc), 3.03 (br, allylic CH), 2.66 (br, allylic CH), 2.08 (br, CHH bridgehead), 2.01(s, OMe), 1.98 (s, OMe), 1.17 (br, CHH bridgehead);  $^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ) 170.9, 170.6, 133.0, 132.2, 76.8, 76.5, 45.3, 40.6, 35.2, 34.3, 33.4, 20.9; IR (nujol)  $1750\text{ cm}^{-1}$  (strong, C=O stretch). DSC analysis reveals a reversible Tg at 102 °C. TGA is shown in Figure 4; isothermal pyrolysis determined the amount loss to be 55.7 % (300 °C, 2 hours). GPC data are found in Table I.

**Typical polymerization of *exo-syn*-2,7-norbornenediacetate (7).** A solution of **7** (431 mg, 2.05 mmol) in THF (10 mL) was added dropwise but quickly to a vigorously stirring solution of Mo(CH-*t*-Bu) (10 mg,  $2. \times 10^{-5}$  mol) in THF (5 mL) over a period of 1-2 minutes. The resulting solution was allowed to stir for 1 hour, quenched with benzaldehyde (15  $\mu$ L, appx 0.15 mmol) and allowed to stir for an additional 30 minutes. The polymer was purified by dropwise addition of the resulting solution into a large excess of methanol, filtration, a subsequent precipitation from chloroform into methanol and placing under vacuum for 24 hours. The polymer was obtained as a white powder (388 mg, 90%):  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  5.2-5.4 (broad multiplets, 2, CHCH), 4.6-4.8 (broad multiplets, 2, CHOAc), 2.6 (br, 1, allylic CH), 2.5 (br, 1, allylic CH), 1.95 (s, 3, OMe), 1.77 (m, 3, OMe);  $^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ) 171.3 and 171.0 (C=O), 133.1 and 132.0 and 131.3 and 130.2 (CHCH, resonances are split due to cis and trans isomers (appx 50:50) and their relation to the acetate group in the 2 position), 79.3, 76.7, 54.2, 46.1, 36.3, 21.3, 21.0; IR (nujol):  $1730\text{ cm}^{-1}$  (strong, C=O stretch). DSC analysis shows a Tg at 110 °C. TGA thermogram is similar to Figure 4 with 56.2 % mass loss observed on heating at 300 °C for 2 hours (calculated for loss of 2 equiv of acetic acid: 57.2 %). GPC data are found in Table I.

**Typical polymerization of exo-cis-2,3-norbornenediol-di-O-isopropylidene (8).** A solution of **8** (344 mg, 2.05 mmol) in THF (5 mL) was added quickly but dropwise to a solution of Mo(CH-*t*-Bu) (10 mg,  $2.05 \times 10^{-5}$  mol) in THF (3 mL) over a period of 1-2 minutes. The color of the solution changed immediately from light yellow to red. The solution was allowed to stir for an additional 30 minutes, quenched with benzaldehyde (15  $\mu$ L) and stirred for an additional 30 minutes to yield a yellow solution. The polymer was obtained free of impurities by precipitation from the above solution into a large excess of methanol followed by three consequent precipitations from chloroform into methanol and removal of solvent in vacuo for 24 hours. The polymer was obtained as a white flaky solid (327 mg, 95%):  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  5.51 (br, trans-CH=CH), 5.28 (br, cis-CHCH), 5.2-5.3 (broad multiplets, CHO), 2.95 (br, allylic CH), 2.55 (br, allylic CH), 2.94 (br, CHH bridgehead), 1.45 (s, CMe), 1.32 (br, CHH bridgehead), 1.23 (s, CMe), integration of resonances consistent with the proposed formulation;  $^{13}\text{C}$  ( $\text{CDCl}_3$ ) 133.0, 131.9, 113.4, 87.1, 86.0, 48.3, 44.3, 38.2, 27.7, 25.2. DSC: T<sub>g</sub> = 102 °C; TGA polymer degradation begins at 300 °C. GPC data are found in Table I.

**Typical base hydrolysis procedure of polyacetate polymers: conversion of poly-6 to poly(exo-syn-2,3-norbornene-diol):** Poly-6(1.0 g) was dissolved in dry THF (150 mL) and added dropwise to a solution of dry THF (150 mL) and 20 mL of a 10% solution of sodium methoxide in methanol under vigorous stirring under reflux. The resulting solution was refluxed for 12 hours, cooled and neutralized with glacial acetic acid. The resulting fine suspension was filtered or centrifuged, washed with methanol and placed under vacuum for 24 hours to give the product as a light yellow powder (0.5 g, 83%) which sparingly soluble in a variety of solvents:  $^1\text{H}$  NMR  $\delta$  ( $\text{CF}_3\text{CO}_2\text{H}:\text{CDCl}_3$ , 1:1) 5.6 (br, 2, CH=CH), 4.21 (br, 1,CHOH), 3.75 (br, 1,CHOH), 2.5-3 (br, 3, OH and allylic CH), 2.05 (br, 3, allylic CH and CH<sub>2</sub>); IR: 3350  $\text{cm}^{-1}$  (br, OH stretch), 1600  $\text{cm}^{-1}$  (C=C stretch) no absorption observed in the carbonyl region.

**Typical acid hydrolysis procedure for poly-8.** Poly-8 (0.3 g) was dissolved in 10 mL of 80% aqueous trifluoroacetic acid and stirred for 3 hours at room temperature. The

resulting solution was cooled to 0°C and slowly neutralized with aqueous KOH (5 M). During the neutralization procedure the color changes from light red to nearly colorless at which time polymer began to precipitate. The resulting product was isolated by filtration or centrifugation, washed several times with water and methanol and placed under vacuum for 24 hours to yield the hydrolyzed polymer as a cream powder (0.2 g, 75%):  $^1\text{H}$  NMR  $\delta$  (MeOD) 5.6 (broad multiplets, 2,  $\text{CH}=\text{CH}$ ), 3.55 (br, 2,  $\text{COH}$  or  $\text{CHOH}$ ), 2.9 (br, 2,  $\text{COH}$  or  $\text{CHOH}$ ), 2.5 (br, 2, allylic  $\text{CH}$ ), 2.0 (br, 1,  $\text{CHH}$ ), 1.1 (br, 1,  $\text{CHH}$ ); IR:  $3350\text{ cm}^{-1}$  (br, OH stretch),  $1590\text{ cm}^{-1}$  ( $\text{C}=\text{C}$  stretch).

**Thermal reaction of acetate polymers.** Thermal treatment of poly-6 and poly-7 was carried out by thermogravimetric analysis (TGA) and temperature programmable pyrolysis oven under a nitrogen atmosphere. Weight loss of the polymers in the powder state was monitored as a function of temperature at a fixed heating rate ( $10\text{ }^\circ\text{C}/\text{min}$ ) or isothermally as a function of time at a given temperature. Polymers in the film state were heated at the same rate and kept isothermally for 2 hours at a given temperature resulting in insoluble black-red films.

**Typical polymerization of 5.** A solution of **5** (122 mg, 1.02 mmol) in THF (1.0 mL) was added quickly to a solution of  $\text{Mo}(\text{CH-}i\text{-Bu})(\text{NAr})(\text{O-}i\text{-Bu})_2$  (10 mg,  $2.05 \times 10^{-5}$  mol) in THF (1.0 mL) and stirred for 15 minutes. Termination was carried out by addition of 15  $\mu\text{L}$  of *p*-trimethylsilyl-benzaldehyde ( $8.4 \times 10^{-5}$  mol). The polymer was purified by precipitation of the polymer from the reaction mixture into 250 ml hexane. Final product is obtained as a white-cream powder; yield 103 mg 85 %:  $^1\text{H}$  NMR ( $\text{CDCl}_3$ )  $\delta$  5.70, 5.55, 5.37, 5.32, 5.20, 3.0, 2.73, 2.55, 2.30, 2.18, 2.03, 1.75, 1.43, (1.004, 0.981, 0.963, 0.956 )( $\text{C}(\text{CH}_3)_3$ ), (0.232, 0.217)( $\text{Si}(\text{CH}_3)_3$ ); the spectrum is too broad for integration within polymer peaks, however integration of *tert*-butyl cap versus olefinic resonance is consistent with assigned stoichiometry as well as 1:1 against TMS cap;  $^{13}\text{C}$  NMR ( $\text{CDCl}_3$ )  $\delta$  135.3, 134.6, 132.3, 131.7, 129.7, 129.2, (CN), 120.9 (CN), 58.5, 44.8, 41.8, 41.2, 39.8, 39.2, 37.0, 36.9, 36.2, 33.8, 29.6 ( $\text{C}(\text{CH}_3)_3$ ), 1.3 ( $\text{Si}(\text{CH}_3)_3$ ).

Mechanical behavior is shown in Figure 7.  $T_g = 123^\circ\text{C}$  by DSC analysis. GPC:  $M_n=24869$  (vs polystyrene),  $\text{PDI}=1.05$ .

mL) was added dropwise but quickly to a light yellow solution of Mo(CH-*t*-Bu) (10 mg,  $2.05 \times 10^{-5}$  mol) in THF (2 mL) under vigorous stirring. After stirring for 15 minutes a solution of **5** (244 mg, 2.05 mmol) in THF (2  $\mu$ L) was added in the same fashion and allowed to stir for 25 minutes to yield a light yellow, slightly viscous solution. The polymerization was quenched by addition of benzaldehyde (15  $\mu$ L) and stirred for an additional 30 minutes. The polymer was isolated by precipitation into a large excess of hexane and centrifugation. The resulting white powder was washed several times with hexane or methanol and placed under vacuum for 24 hours to yield 381 mg (87 %) of the product. The  $^1\text{H}$  and  $^{13}\text{C}$  NMR spectra were identical to the superposition of spectra for poly-NBE and poly-**5**. GPC:  $M_n$ =55780 (vs polystyrene), PDI=1.05. Mechanical behavior is shown in Figure 7.

**Acknowledgements.** This work was supported in part by the Office of Naval Research. G.B. thanks NSERCC for a predoctoral Fellowship. We also thank Dr. Vernon Gibson and Prof. James W. Feast for samples of **9**, **10** and **11** and carbonate.

---

## References

1. (a) Ivin, K. J. *Olefin Metathesis*, Academic Press: London, 1983. (b) Dragutan, V.; Balaban, A. T.; Dimonie, M. *Olefin Metathesis and Ring-opening Polymerization of Cyclo-Olefins*, 2nd Ed., Wiley-Interscience, 1985.
2. Schrock, R. R.; Murdzek, J. S.; Bazan, G. C.; Robbins, J.; DiMare, M.; O'Regan, M. J. *Am. Chem. Soc.* **1990**, *112*, 3875.
3. Schrock, R. R.; DePue, R. T.; Feldman, J.; Yap, K. B.; Yang, D. C.; Davis, W. M.; Park, L. Y.; DiMare, M.; Schofield, M.; Anhaus, J.; Walborsky, E.; Evitt, E.; Krüger, C.; Betz, P. *Organometallics* **1990**, *9*, 2262.
4. Schrock, R. R. *Acc. Chem. Res.* **1990**, *24*, 158.
5. Murdzek, J. S.; Schrock, R. R. *Macromolecules* **1987**, *20*, 2640.
6. (a) Bazan, G.; Khosravi, E.; Schrock, R. R.; Feast, W. J.; Gibson, V. C. *Polymer*



---

*Commun.* **1989**, *30*, 258. (b) Bazan, G.; Khosravi, E.; Schrock, R. R.; Feast, W. J.; Gibson, V. C.; O'Regan, M. B.; Thomas, J. K.; Davis, W. M. *J. Am. Chem. Soc.*, in press.

7. Crowe, W. E.; Mitchell, J. P.; Gibson, V. C.; Schrock, R. R. *Macromolecules* **1990**, *23*, 3534.

8. (a) Gilliom, L. R.; Grubbs, R. H. *J. Am. Chem. Soc.* **1986**, *108*, 733. (b) Grubbs, R. H.; Tumas, W. *Science* **1989**, *243*, 907.

9. Bazan, G. C.; Schrock, R. R. *Macromolecules*, in press.

10. (a) Gold, L. *J. Chem. Phys.* **1958**, *28*, 91. (b) Rempp, P.; Merrill, E. W. *Polymer Synthesis*. New York: Huethig and Wepf; 1986.

11. Novak, B. M.; Grubbs, R. H. *J. Am. Chem. Soc.* **1988**, *110*, 7542.

12. Bazan, G. C.; Schrock, R. R.; O'Regan, M. B. *Organometallics*, in press.

13. Oskam, J.; unpublished observations.

14. Matsumoto, S.; Komatsu, K.; Igarashi, K. *Polymer Preprints*, **1977**, *18(1)*, 110.

15. Feast, W. J.; Harper, K. *Brit. Polym. J.* **1986**, *18*, 161.

16. Feast, W. J.; Harper, K. *J. Mol. Cat.* **1985**, *28*, 293.

17. (a) Feldman, J.; Davis, W. M.; Thomas, J. K.; Schrock, R. R. *Organometallics* **1990**, *9*, 2535. (b) Feldman, J.; Murdzek, J. S.; Davis, W. M.; Schrock, R. R. *Organometallics* **1989**, *8*, 2260.

18. Kelen, T. *Polymer Degradation*, van Nostrand Reinhold, New York, 1981.

19. Gardner, D. L.; McNeill, I. C. *J. Thermal Anal.* **1969**, *1*, 389.



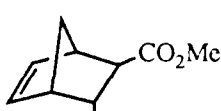
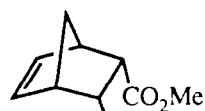

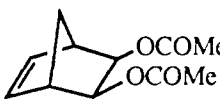
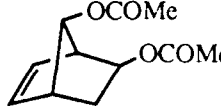
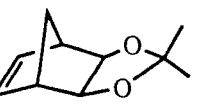
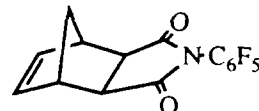


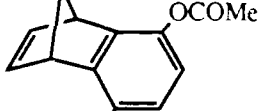
20. (a) Wypych, J. *Polyvinyl Chloride Degradation*, Elsevier, Amsterdam, 1985. (b) Geddes, W. C. *Eur. Polym. J.* **1967**, *3*, 474.

21. (a) Finch, C. A. *Poly Vinyl Alcohol*, John Wiley & Sons, New York, 1973. (b) Maruyama, K.; Yagi, M.; Tanizaki, Y. *Polymer Commun.* **1986**, *27*, 349. (c) Maruyama, K.; Take, M.; Fujii, N.; Tanizaki, Y. *Bull. Chem. Soc. Jpn.* **1986**, *59*, 13. (d) Maruyama, K.; Kuramoto, Y.; Tanizaki, Y. *Polymer* **1989**, *30*, 1419.

22. Ramakrishnan, S.; Chung, T. C. *Macromolecules* **1989**, *22*, 3188.

- 
23. Craig, D. *J. Am. Chem. Soc.* **1951**, *73*, 4889.
  24. Castner, K. F.; Calderon, N *J. Mol. Cat.* **1982**, *16*, 47.
  25. Brace, N. O. *J. Org. Chem.* **1979**, *44*, 1964.
  26. Shealy, Y.; Clayton, J. D. *J. Am. Chem. Soc.* **1969**, *91*, 3075.
  27. Lambert, J. B.; Holcomb, A. G.; *J. Am. Chem. Soc.* **1971**, *93*, 2994.
  28. Just, G.; Reader, G.; Chalard-Faure, B. *Can. J. Chem.* **1976**, *54*, 849.

Table I. Monomers polymerized by  $\text{Mo}(\text{CH-}t\text{-Bu})(\text{NAr})(\text{O-}t\text{-Bu})_2$ .

 <b>1</b>	 <b>2</b>	 <b>3</b>	 <b>4</b>
50 <sup>a</sup> /1.07 <sup>b</sup> 100/1.06 200/1.04	50/1.06 100/1.07	50/1.05 100/1.04 200/1.04	50/1.11 100/1.12 200/1.22
 <b>5</b>	 <b>6</b>	 <b>7</b>	 <b>8</b>
50/1.06 100/1.07 200/1.07	50/1.14 100/1.19	50/1.13 100/1.11	50/1.12 100/1.07
 <b>9</b>	 <b>10</b>	 <b>11</b>	 <b>12</b>
50/1.05	100/1.07	100/1.06 <i>+ block</i>	100/1.16 <sup>c</sup>

<sup>a</sup>Equivalents of monomer added. <sup>b</sup>Polydispersity index (vs. polystyrene in dichloromethane). <sup>c</sup>Bimodal GPC trace. For reaction conditions refer to experimental section.

Table II. Distribution of products as a function of temperature after dissolving **14** in toluene-d<sub>8</sub>.

Temp (°C)	Mo(CHCMe <sub>3</sub> )	<b>14</b>	<b>15</b>	FIP <sup>a</sup>	SIP <sup>b</sup>
-45	1	4	1	trace	
0	1	2	0.9	0.2	
25	1	0.1	0.9	0.3	0.06

<sup>a</sup>Alkylidene due to first insertion product. <sup>b</sup>Second insertion product. Equilibration time at each temperature was 20 min.

## Figure Captions.

**Figure 1.** 500 MHz  $^1\text{H}$  NMR spectrum ( $\text{CDCl}_3$ ,  $40^\circ\text{C}$ ) in the olefinic region of the organic products resulting from adding 20 eq of **2** to  $\text{Mo}(\text{CH-}t\text{-Bu})(\text{NAr})(\text{O-}t\text{-Bu})_2$  followed by a benzaldehyde quench. (\* = *trans*- $\text{PhCH=CH-}t\text{-Bu}$ )

**Figure 2.** 500 MHz  $^1\text{H}$  NMR spectrum ( $\text{C}_6\text{D}_6$ ) in the alkylidene region of a living 20 mer of poly-**3**.

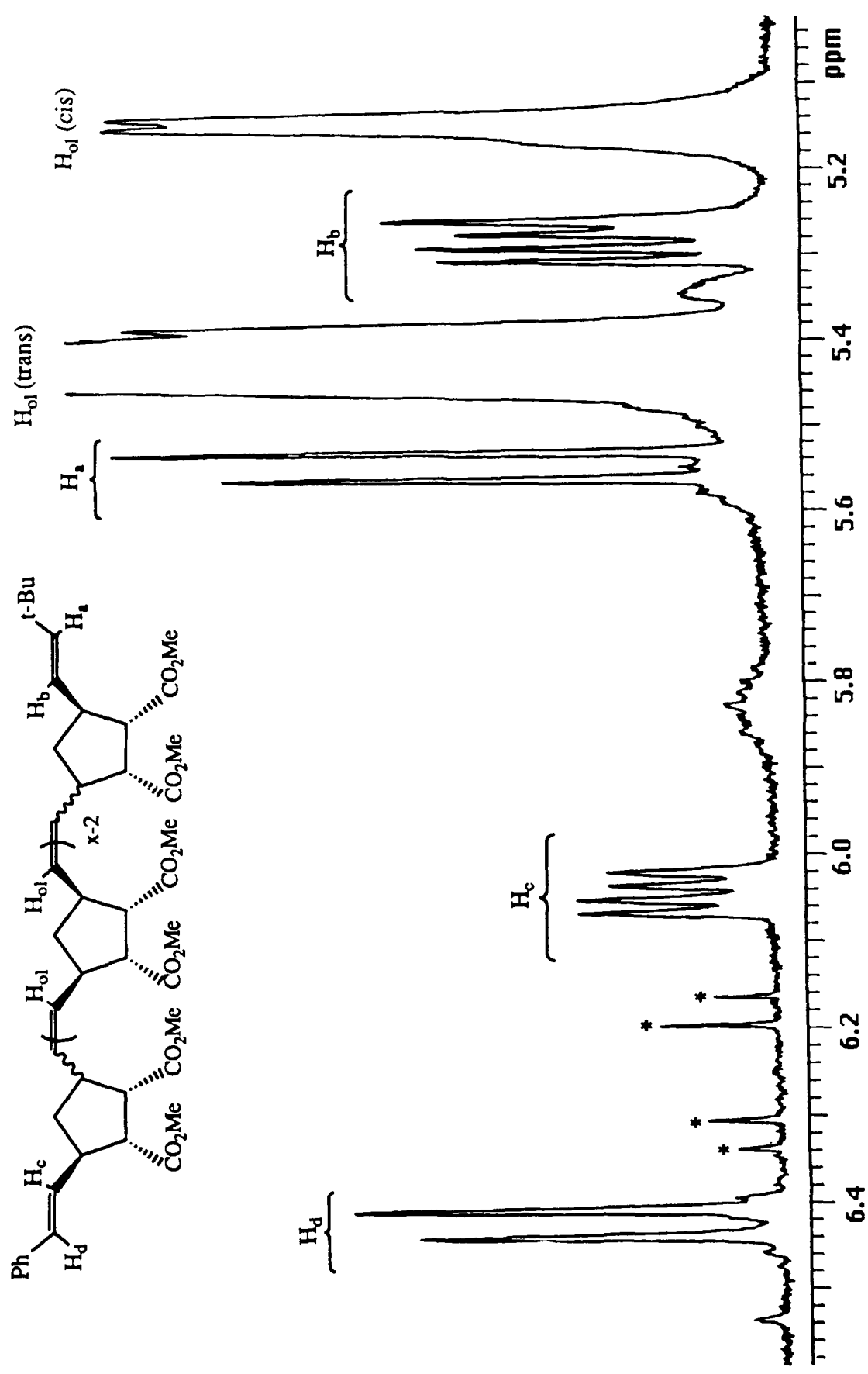
**Figure 3.** 500 MHz  $^1\text{H}$  NMR spectrum ( $\text{THF-}d_8$ ) in the alkylidene region of a living 20 mer of poly-**5**.

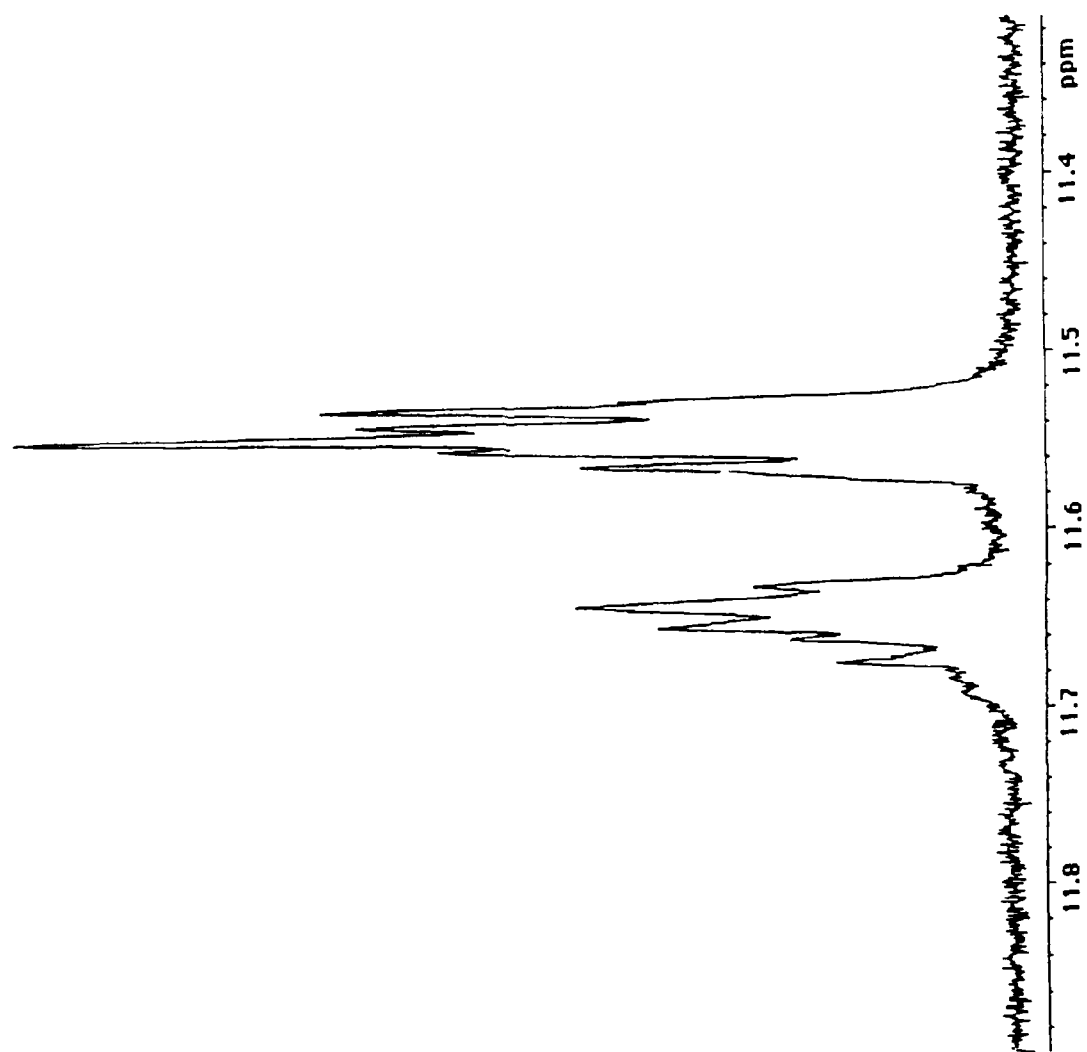
**Figure 4.** TGA analysis of poly-**6**.

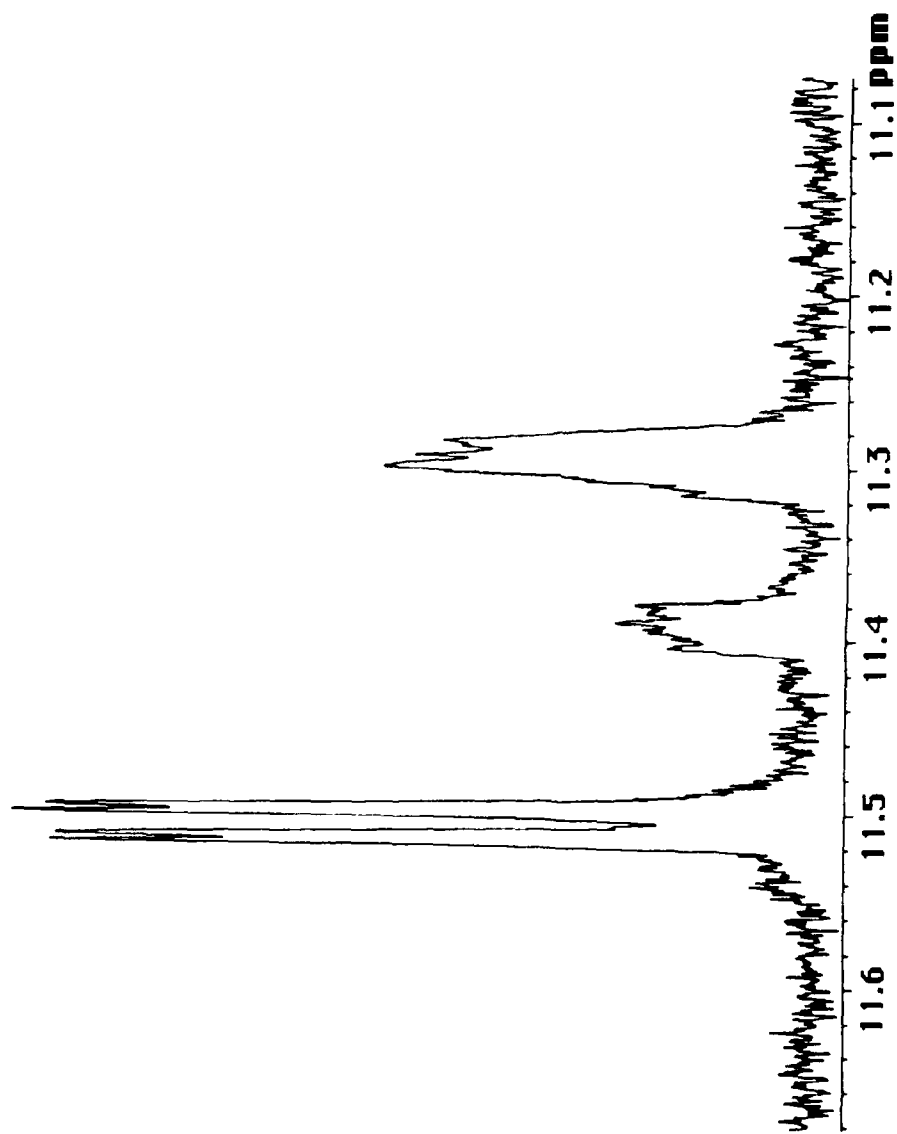
**Figure 5.** GPC trace of poly-**1/5**.

**Figure 6.** 125 MHz  $^{13}\text{C}$  NMR ( $\text{CDCl}_3$ ) spectra in the olefinic region of (a) poly-**5**; (b) poly-**1/5**.

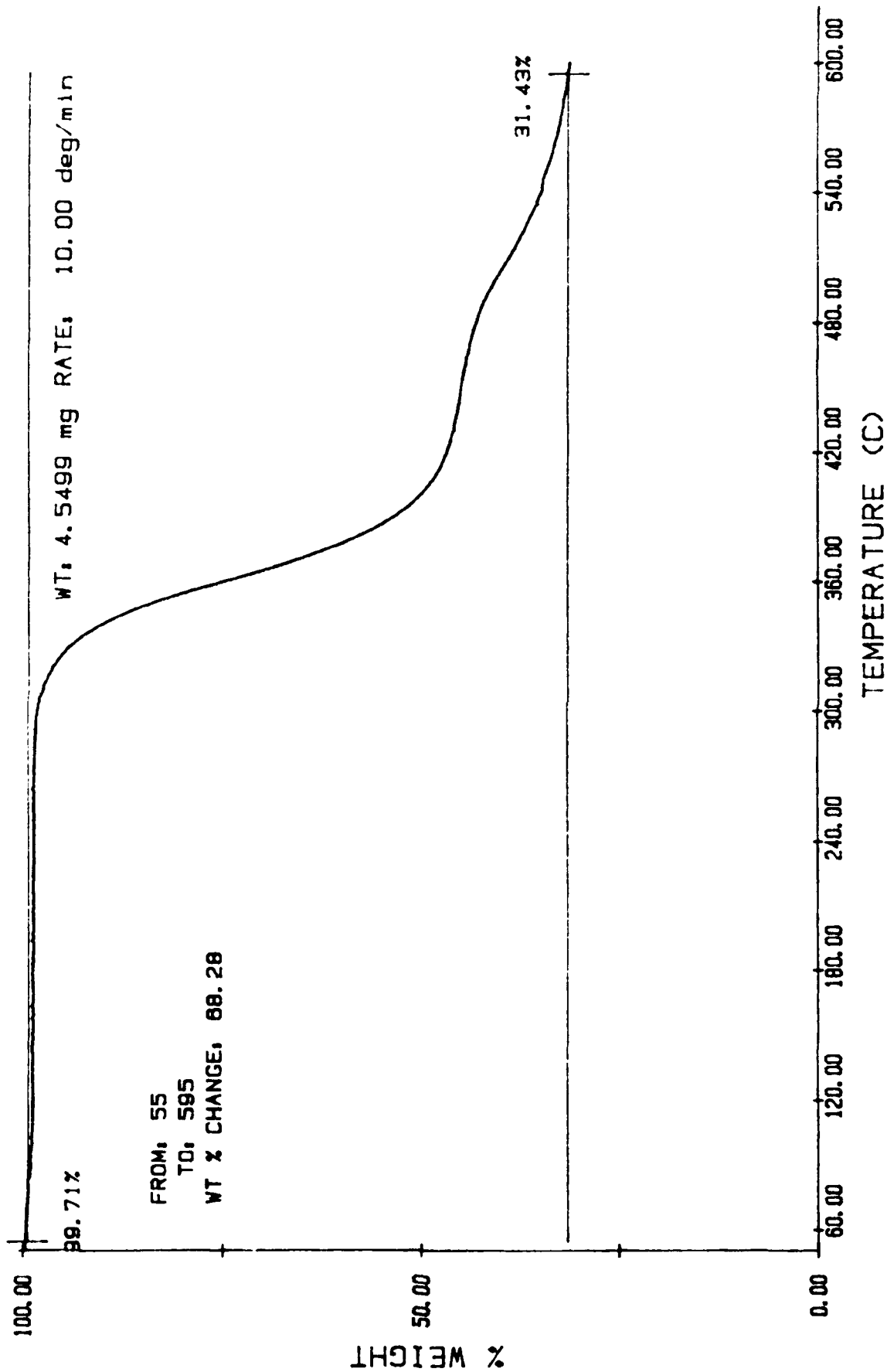
**Figure 7.** Stress-strain curves (10 mm/min) of (a) poly-**5**; (b) poly-**1/5**.



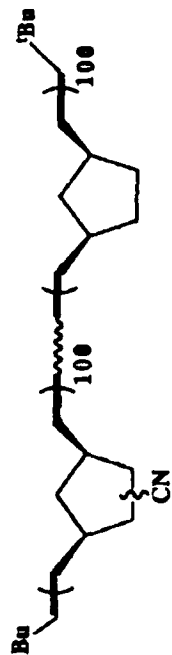






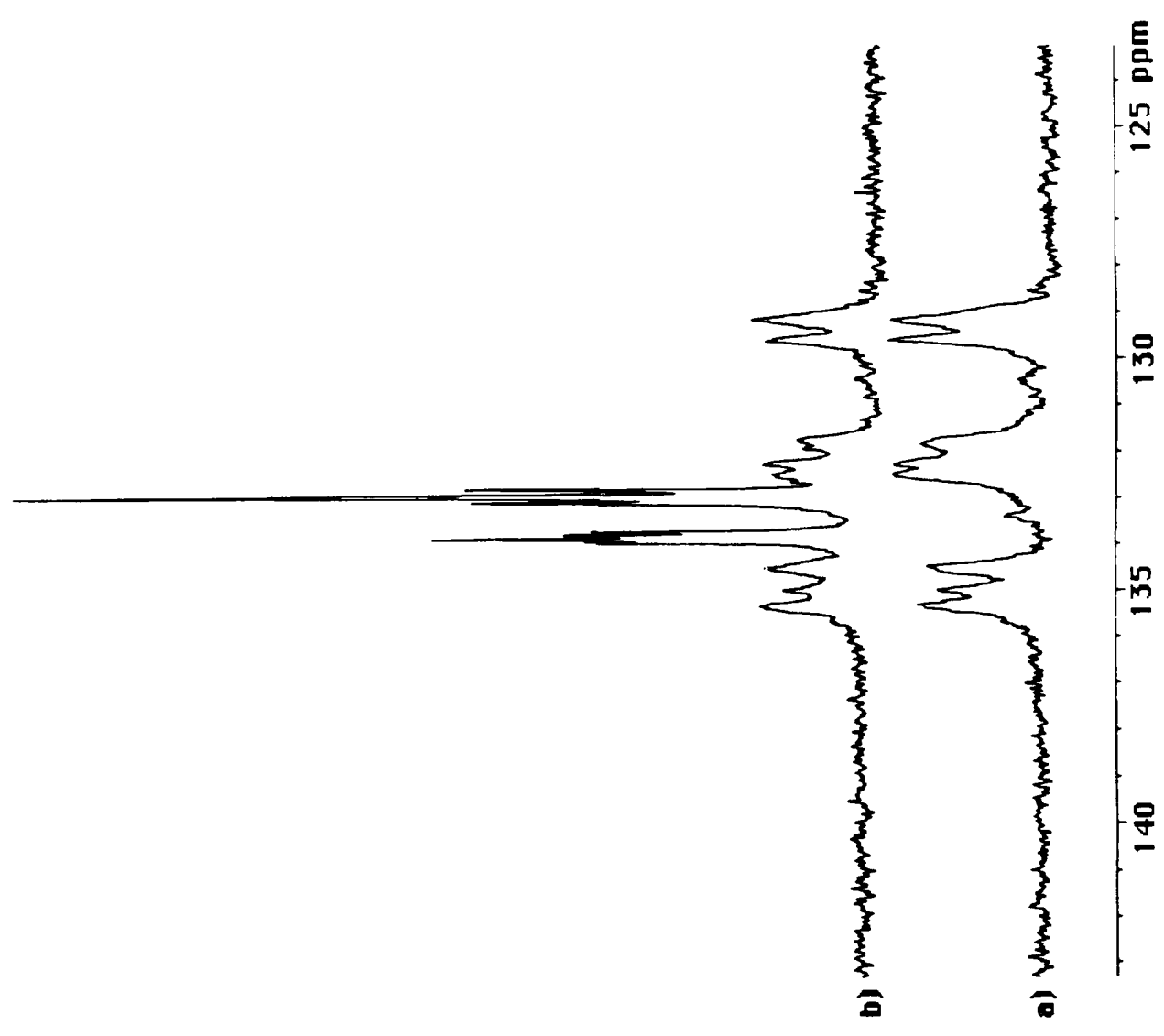


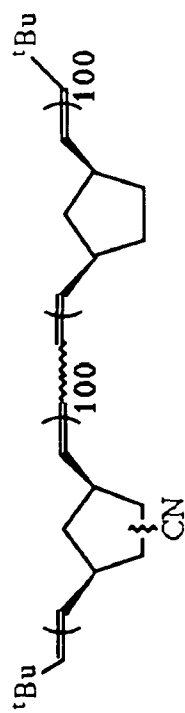
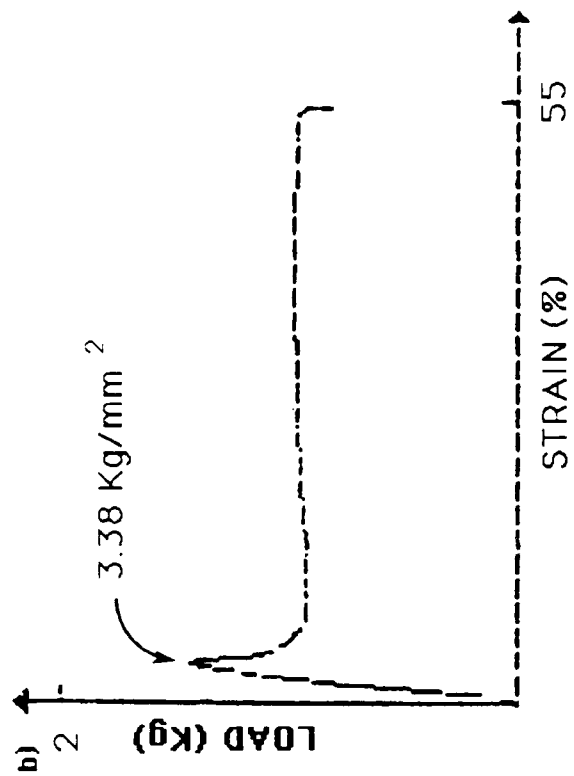
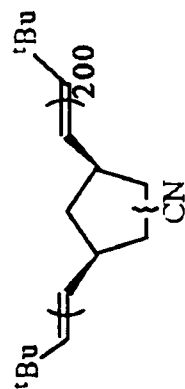
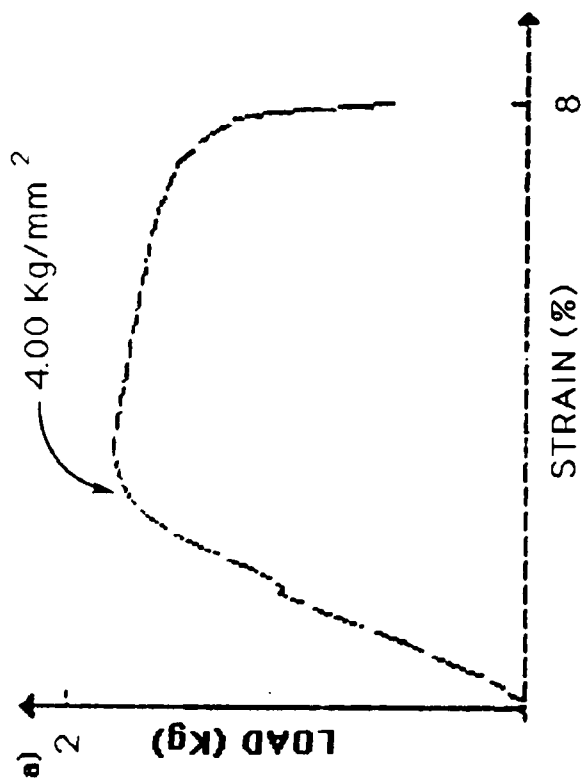
REFRACTOMETER DATA



PDI = 1.05

10 14.00 18.00 22.00 26.00 30.00Min





LIST 5

Dr. Harry R. Allcock  
Department of Chemistry  
Pennsylvania State Univ.  
University Park, PA 16802

4132007

Dr. Chris W. Allen  
Department of Chemistry  
University of Vermont  
Burlington, VT 05405

413c012

Dr. J. M. Augl  
Naval Surface Weapons Center  
White Oak, MD 20910

Dr. Kurt Baum  
Fluorochem, Inc.  
680 S. Ayon Avenue  
Azusa, CA 91702

4000021sbi

Dr. Len J. Buckley  
Naval Air Development Center  
Code 6063  
Warminster, PA 18974

Dr. Ivan Caplan  
DTNSRDC Annapolis  
Code 0125  
Annapolis, MD 21401

Dr. E. Fischer  
DTNSRDC Code 2853  
Annapolis, MD 21402

Dr. Robert H. Grubbs  
Department of Chemistry  
California Inst. of Technol.  
Pasadena, CA 91124

4132019

Dr. Henry K. Hall  
Department of Chemistry  
University of Arizona  
Tucson, AZ 85721

413j009

Dr. T. J. Reinhart, Jr.  
Nonmetallic Materials Div.  
AF Materials Lab. (AFSC)  
Wright-Patterson AFB,  
OH 45433

Dr. Richard M. Laine  
Washington Technology Center  
University of Washington  
Seattle, WA 98195

s400033srh

Dr. Robert W. Lenz  
Polymer Sci. and Eng. Dept.  
University of Massachusetts  
Amherst, MA 01002

441c013

Dr. Krzysztof Matyjaszewski  
Department of Chemistry  
Carnegie-Mellon University  
Pittsburgh, PA 15213

413j002

Dr. Dietmar Seyferth  
Department of Chemistry  
Massachusetts Inst. of Techn.  
Cambridge, MA 02139

413c004

Dr. James E. McGrath  
Department of Chemistry  
Virginia Polytechnic Inst.  
Blacksburg, VA 24061

4132007

Dr. L. E. Slotter  
Code Air 931-A  
Naval Air Systems Command  
Washington, D. C. 20361-9310

Dr. William B. Moniz  
Code 6120  
Naval Research Laboratory  
Washington, DC 20375-5000

4132026

Dr. James M. Tour  
Dept. of Chemistry  
Univ. of South Carolina  
Columbia, SC 29208

400x056

Dr. James A. Moore  
Department of Chemistry  
Rensselaer Polytechnic Inst.  
Troy, NY 12180-3590

413c014

Dr. David M. Walba  
Dept. of Chem. & Biochem.  
Univ. of Colorado  
Boulder, CO 80309

413h010

Dr. Virgil Percec  
Dept. of Macromolecular Sci.  
Case Western Reserve Univ.  
Cleveland, OH 44106-2699

413c024

Dr. Richard R. Schrock  
Department of Chemistry  
Massachusetts Inst. of Techn.  
Cambridge, MA 02139

4132038